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Result
No.
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Maximum DB seq length: 2000000000
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Listing first 45 summaries
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7: /net/abss06/SIDS1/gcgdata/hold-geneseq/geneseqn-embl/NA1986.DAT:*
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Human kinase PKIN-	7	for	3	Human brain expres	Probe #7320 for ge	$\vdash$	Drosophila melanog	Human POLY3 cDNA.	Human POLY2 cDNA.	Novel protein kina	Human POLY4 cDNA.	Trad	n Trad	ID No:	ID No: 28	Novel protein kina		Novel protein kina	Nucleotide sequenc	Polynucleotide seq	Drosophila melanog	Drosophila melanog	Drosophila melanog	Human cDNA clone (	Novel protein kina	cDNA encoding huma		æ	Human prostate can	Human bone marrow	Human bone marrow	Human immune/haema	Human cDNA clone r	an cDNA 5'	DNA sequence encod

## ALIGNMENTS

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RESULT
AAZ49765
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15-JUN-1998;
                                          15-JUN-1999;
                                                                                     23-DEC-1999
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98US-0089294
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Best Local Similarity
Matches 789; Conser
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kinase-related protein 1 (DRP-1), which is a calmodulin-dependent
serine/threonine kinase. DRP-1 is a cytoplasmic protein capable of
inducing apoptosis by dimerisation. It shows significant homology to
DAP kinase. It has cytostatic, antipsoriatic and immunosuppressive
activity and can be used for inhibiting growth/metastasis of
tumours and promoting death of tumour cells. It can also be used in
the treatment of cancer, psoriasis and autoimmune diseases. Fragments
of DRP-1 DNA are useful as probes for screening individuals with a
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MCINNIS P A.
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cc polymerase chain reaction (PCR) primers, oligomers, and for chromosome cc polymerase chain reaction (PCR) primers, oligomers, and for chromosome cc polymcleotides are also used in diagnostics as expressed sequence tags cfor identifying expressed genes. (I) is useful in gene therapy techniques cc for identifying expressed genes. (I) is useful in gene therapy techniques cc (II). (II) is useful for generating antibodies against it, detecting or quantitating a polypeptide in tissue, as molecular weight markers and as a food supplement. (II) and its binding partners are useful in medical cc imaging of sites expressing (II). (I) and (II) are useful in medical cc imaging of sites expressing (II). (I) and (II) are useful for treating cc diagnostics, forensics, gene mapping, identification of mutations cresponsible for genetic disorders or other traits to assess biodiversity and no acid sequences. AAS64197-AAS94564 represent novel human cc diagnostic coding sequences of the invention.

Cc specification, but was obtained in electronic format directly from WIPO
                                                                                                                                                                                                                                                                                                                         New isolated polynucleotide and encoded polypeptides, useful in diagnostics, forensics, gene mapping, identification of mutations responsible for genetic disorders or other traits and to assess biodiversity -
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23-AUG-2000;
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P-PSDB; ABG09274.
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length cDNA; cDNA synthesis; oligo-capping;
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11-JAN-2000;
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                                                                                                                                                                                                                                                                                                                                                            molecules have been determined. Primers for synthesising the full length cDNA are useful for clarifying the function of the protein encoded by the cDNA. The full length clones were obtained by construction of full length enriched cDNA libraries that were synthesised by the oligo-capping method. The primers enable the production of the full length cDNA easily without any special methods. The present sequence is a full length human cDNA of the invention.

Note: The sequence data for this patent did not form part of the printed specification, but was obtained in CD-ROM format directly from EPO.
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The invention provides human and murine recombinant Zipper Interacting Protein Kinase (ZIP-kinase) proteins. These proteins are serine/threonine kinases which bind the leucine zipper domain of transcription factor ATF4. Host cells containing vectors comprising the ZIP-kinase nucleic acids are used for the recombinant expression of the proteins. ZIP-kinase protein and DNA are useful as gene therapeutic agents against cancer, and as anti-cancer agents. The present sequence represents a DNA encoding a
                                                                                                                      New Recombinant Zipper Interacting Protein Kinase protein and DNA, useful as anticancer agents
                                                                                                  Claim 6; Page 19-22;
                                                                                                                                                                                                 Akira
                                                                                                                                                                                                                       (NISC-)
                                                                                                                                                                                                                                             26-SEP-1997;
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Human

ZIP-kinase

(serine/threonine kinase)

encoding

DNA.

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Best Local Similarity 76.6%;
Matches 604; Conservative
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tein and DNA, useful as anticancer
                     aagaaaattgctcactttgatctcaagccagaaaacattatgttgttagacaagaatatt
                                                                                                           cttgagctagtgtctggaggagagctcttcgatttccttggcccagaaggagtcactgagt
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ne zipper domain;
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27-AUG-1999;
11-JAN-2000;
02-MAY-2000;
09-JUN-2000;
                                                                                                                                            Primer sets for synthesizing polynucleotides, particularly the full-length cDNAs defined in the specification, and for the detand/or diagnosis of the abnormality of the proteins encoded by full-length cDNAs -
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Human cDNA
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99JP-0300253.
2000JP-0118776.
2000JP-0183767.
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A, Nagai K,
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The present invention describes primer sets for synthesising 5602 full-length cDNAs defined in the specification. Where a primer set comprises: (a) an oligo-dT primer and an oligonucleotide complementary to the complementary strand of a polynucleotide which comprises one of

8;

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ROM;

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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       sequence and an oligonucleotide comprising a sequence complementary to a polynucleotide which comprises a 3'-end sequence, where the oligonucleotide which comprises a 1-end sequence, where the street sequence of the 5'-end sequence and the combination of the 5'-end sequence and sequence is selected from those defined in the specification. The primer sets can be used in antisense therapy and in gene therapy. The primers are useful for synthesising polynucleotides, particularly full-length cDNAs. The primers are also useful for the detection and/or diagnosis of the abnormality of the proteins encoded by the full-length cDNAs. The primers allow obtaining of the full-length cDNAs easily without any specialised methods. AAH03166 to AAH13628 and AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to AAH3630 represent oligonucleotides, all of which are used in the exemplification of the present invention.
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970

Query Match Best Local S Matches 604

Similarity

62.6%; 76.6%;

Conservative

0

184;

Indels Length

0;

Gaps

0,

Score 493.6; DB Pred. No. 6.4e-1 0; Mismatches 1

DB

22;

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. Local Simes 604;

Sequence

2224

BP;

419 A;

656 C;

806 G;

343

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                                   phosphatase. The polypeptides are expected to participate in signal transduction in cells. The kinase phosphatases are connected with intracellular signaling pathways. Antisense oligonuclectides and compounds identified by screening (agonists or antagonists) can be used to treat human or animal disorders associated with the expression or function of the protein. In addition, the polypeptides may be used as target molecules fir drug development.
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Ishii
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11-JAN-2000;
17-FEB-2000;
                                                                                                                                                                                                                                                             Claim 1; Page 119-125; 336pp; Japanese
                                                                                                                                                                                                                                                                                                        New genes encoding protein kinase and protein phosphatase, use identifying modulators which can be used to treat human or and disorders associated with the expression or function of these
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T, Wakamatsu
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                                                                                                                                associated
                                                                                                                                                                                death
                                                                                                                                                                           associated protein
                                                                                                                                                                                                                                (first entry
Location/Qualifiers
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                                                                                                                                                                                                                                                                                                                                      CDNA; 4272
                                                                                                                             DAP;
                                                                                                                                                                                                                                                                                                                                      ВP
                                                                                                                          cytokine;
                                                                                                                                                                             DAP-2.
                                                                                                                             cell
                                                                                                                             death;
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aaaagtaccggcctccagtatcccgccaaattcatcaagaaaaggaggactaagtccagc aagagcacggggcttgagtatgcagccaagttcatcaagaagcggcagagccgggcgagc 120 tacgacaccggcgaggaacttggcagtggacagtttgcggttgtgaagaaatgccgtgag

156

Query Match Best Local Sim Matches 557;

Similarity

50.5%;

Score 398.2; DB 16 Pred. No. 1.8e-103; Mismatches

DB 16;

4272; 24;

Indels Length

Gaps

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96

Conservative

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tatgacatcggagaggagctgggggagtggccagtttgccatcgtgaagaagtgccgggag 60

Sequence 4272 BP; 1076 A; 1161 C; 1121 G; 914 T; 0

other

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cc was cloned in antisense orientation into the EBY-based pTKO1 cc expression vector. The resulting expression library was introduced into HeLa cells. A fraction of the transfectants was selected with CC hydromycin B. The majority of transfected cells were selected with CC both hydromycin B and IFN-gamma. The cells that survived and/or CC grew in the presence of IFN-gamma were expanded and pooled. The CC which included DNA was obtd. and cleaved with Dpn1 and introduced CC which included DNA antisense sequences, some of which were able to CC protect cells from the death-promoting effects of IFN-gamma. CC plasmid DNAs were prepd. from 10 individual bacterial clones. PCR CC primers that corresp. to the immediate flanking sequence of the CC DNA insertion sites in the pTKO1 vector. The PCR fragments were CC used as labeled probes to seach Southern blots for possible cross CC hybridisation between some of the rescued antisense cDNA clones were classified into six distinct CC non-overlapping yps., some constituting several members (clones) CC and some constituting a single member. Antisense cDNA clones CC the DNA product called DAP-2. Clone 256 (DAP-2) was sequenced and CC composite sequence derived from 2 clones and the deduced AA sequence CC canson in AA089838 and AAR74205. The ORF is also shown in AA089839. CC and shown in AA089839 and AAR74205. The ORF is also shown in AA089839. CC in the protein (see AAQ74205 FT).
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              DNA whose expression mediates cytokine-induced programmed cedeath - used to treat diseases or disorders associated with uncontrolled, pathological cell growth or cytokine-induced
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            WPI;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         DAP genes seem to play an imp. role in programmed cell death and the inhibition of their expression protects the cell from cytokine-promoted cell death. A cDNA library was generated from a mixture of mRNAs harvested after treatment of HeLa cells with IFN-gamma. It
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Claim 2; Fig 8; 61pp;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            programmed cell death.
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(YEDA ) YEDA F
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В γ В ρy

121 433

cggcgcggtgtgagccgggaggatcgagcgggaggtgagcatcctgcggcaggtgctg aaaagtaccggcctccagtatcccgccaaattcatcaagaaaaaggaggactaagtccagc aagagcacggggcttgagtatgcagccaagttcatcaagaagcggcagagccgggcggagc tacgacaccggcgaggaacttggcagtggacagtttgcggttgtgaagaaatgccgtgag Matches

Conservative

0,

Score 398.2; DB 16 Pred. No. 2.1e-103; 0; Mismatches 208;

Indels Length

24;

Gaps

1;

432 60 DB 16;

1500 G; 1415 T; 0 other;

373 61

1 tatgacatcggagaggagctggggagttggccagtttgccatcgtgaagaagtgccgggag

Query Match Best Local Similarity

50.5%; 70.6%;

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was cloned in antisense orientation into the EBV-based pTKMU
expression vector. The resulting expression library was introduced
into HeLa cells. A fraction of the transfectants was selected with
lygromycin B. The majority of transfected cells were selected with
both hygromycin B and IFN-gamma. The cells that survived and/or
grew in the presence of IFN-gamma were expanded and pooled. The
extrachromosomal DNA was obtd. and cleaved with Dpnr and introduced
into E. coli HB101 host cells. A few bacterial clones were obtd.
which included DNA antisense sequences, some of Which were able to
protect cells from the death-promoting effects of IFN-gamma.
Plasmid DNAs were prepd. from 10 individual bacterial clones. PCR
amplified CDNA inserts were generated from each plasmid using
primers that corresp. to the immediate flanking sequence of the
CDNA insertion sites in the pTK01 vector. The PCR fragments were
hybridisation between some of the rescued antisense cDNA clones.
                                 The 10 cDNA clones were classified into six distinct non-overlapping gps., some constituting several members (clones) and some constituting a single member. Antisense cDNA clone 256 has the DNA product called DAP-2. Clone 256 (DAP-2) was sequenced and used to screen a K562 lambda gt10 cDNA library. The resulting composite sequence derived from 2 clones and the deduced AA sequence are shown in AAQ89338 and AAR74205. The ORF is also shown in AAQ89339. AAQ89388 has a poly A tail. The calculated mol. wt. of the protein is about 160 kDa. Several known domains and motifs were identified in the protein (see AAQ74205 FT).
Sequence 5886 BP;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       DAP genes seem to play an imp. role in programmed cell death and the inhibition of their expression protects the cell from cytokine-promoted cell death. A cDNA library was generated from a mixture of mRNAs harvested after treatment of HeLa cells with IFN gamma. It
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         DNA whose expression mediates cytokine-induced programmed death - used to treat diseases or disorders associated wi uncontrolled, pathological cell growth or cytokine-induced programmed cell death.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Kimchi A;
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/label= instability
1447 A; 1524 C;
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RESULT 1
AAVO2289
ID AAVV
XX AVV
XX DAVV
XX DEAT
WWW METER
KW METER
KW METER
KW ALZI
XX DEAT
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                                                                                                                                    metastatic activity; cancer; psoriasis; autoimmune disea; programmed cell death; degenerative neurological disease
                                                                                                                                                                                  DNA sequence encoding
  11-SEP-1998
                       WO9839429-A2
                                                                                                                           Alzheimer's;
                                                                                                                                                            Death associated protein; DAP-2; cell death; tumour cell; DAP-kinase
                                                                                                                                                                                                           12-JAN-1999
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                                                                                                                                                                                                                                                      standard;
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                                             /*tag= a
/product=
                                                                   Location/Qualifiers 337..4608
                                                                                                                                                                                                                                                      DNA;
                                                                                                                                                                                                         entry)
                                                                                                                                                                                 death associated protein-2 (DAP-1, DAP-kinase).
                                              DAP-2
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       The present sequence encodes a death associated protein-2 (DAP-2, DAP-kinase). The DAP genes and proteins are used for promoting death of normal or tumour cells, and for suppressing the metastatic activity of tumour cells. They can be used in the treatment of diseases or disorders associated with uncontrolled pathological growth, e.g. cancer, psoriasis, autoimmune diseases and others. Agents which antagonise, inhibit or nuetralize DAP products are used for protecting cells from programmed cell death. In this case they can be used for the treatment of degenerative neurological diseases, e.g. Alzheimer's, prevention of death of T cells in AIDS patients, prevention of rejection associated cell death in transplants, and protection of normal cells from the cytotoxic effects of anti-cancer therapies.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      New isolated death associated protein nucleic acids - used for the diagnosis and treatment or disorders associated with programmed cedeath, e.g. cancers, auto:immune disease or neurological disease
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P-PSDB; AAW71367.
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                                                                                                                                      aagaaaattgctcactttgatctcaagccagaaaacattatgttgttagacaagaatatt 420
gaatttaagaatatttttgggacgccggaatttgttgctccagaaattgtgaactacgag
                                                                   cccattccacacatcaagctgattgactttggtctggctcacgaaatagaagatggagtt
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                                                   cccaaacctcggatcaagatcattgactt--
                                                                                                                    cttcaaatcgcccactttgatcttaagcctgagaacataatgcttttggatagaaatgtc
                                                                                                                                                                                        gaagaggaagcaactgaatttctcaaacaaattcttaatggtgtttactacctgcactcc
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Pred. No. 2.1e-103;
0; Mismatches 208;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              0 other
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540 828 480 732 360 672 300 612 240

792

552 180 120 432 60

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B 64

46.5%; 75.9%;

Score 366.6; DB 22; Pred. No. 8.7e-95;

Length

757;

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RESULT 11
AAK9185
XX AAK918
XX AAK918
XX D6-NOV
XX Human;
XX Human;
XX Ep1130
XX Ep113
                            The invention relates to primers for synthesising full length cDNA clones. 830 cDNA molecules encoding a human protein have been isolated and nucleotide sequences of 5'- and 3'-ends of the cDNA molecules have been determined. Primers for synthesising the full length cDNA are useful for clarifying the function of the protein encoded by the cDNA. The full length clones were obtained by construction of full length enriched cDNA libraries that were synthesised by the oligo-capping without any special methods. The production of the full length cDNA easily sequence of the 5'-end of a cDNA provided in the invention.

Note: The sequence data for this patent did not form part of the printed specification, but was obtained in CD-ROM format directly from EPO.
                                                                                                                                                                                                                                                                                                                                                                                                    Claim 2; SEQ ID NO 316; 1380pp + sequence listing; English
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11-JAN-2000; 2000JP-0118774
02-MAY-2000; 2000JP-0183765
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Nagai
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S, Otsuki
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T, Koga
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9 other;

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Matches 486; Conservative
  08-JUL-1999;
11-JAN-2000;
02-MAY-2000;
                                              07-JUL-2000;
                                                                      05-SEP-2001.
                                                                                            EP1130094-A2
                                                                                                                                                            Human cDNA clone representative sequence,
                                                                                                                                           Human; full length
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                                                                                                                                                                                                                                   AAK93262 standard; cDNA; 757
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99JP-0194486.
2000JP-0118774.
2000JP-0183765.
                                               2000EP-0114089
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Note: The sequence data for this patent did not form part of the printed specification, but was obtained in CD-ROM format directly from EPO.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           WPI; 2001-524255/58.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Example 11; SEQ ID NO 1722;
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Wakamatsu A, Sugiyama
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               accgctgggccttggaagcngacatgttgaacatccgtgtctcccctatatcctcctgaa
                       gcccctgggtc-tggaggctgacatgtggagcataggcgtcatcacctacatcctcttaa
                                                                                              gaattcaanaacatcttcggcaccccggaattttgtggccccanaaattgtgaactatga
                                                                                                                     gaatttaagaatatttttgggacgccggaa-tttgttgctcccagaaattgtgaactacga
                                                                                                                                                                                                 cccaaccacgaatcaagctcatcgacttcggcatcgcncacaagatcgangcggggaac
                                                                                                                                                                                                                        cccattccacacatcaagctgattgactttggtctggctcacgaaatagaagatggagtt
                                                                                                                                                                                                                                                                                                 aagcgcatcgcacactttgacctgaagccggaaaacatcatgctgctggacaagaacgtg
                                                                                                                                                                                                                                                                                                                         aagaaaattgctcactttgatctcaagccagaaaacattatgttgttagacaagaatatt 420
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     ctggagctggtctctggcggggagctctttgacttcctggcggagaaggagtcgctgacg
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m T, Nagai
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K, Kojima
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2000US-0186350. 2000US-0189874. 2000US-0190076. 2000US-0198123. 2000US-0205515.

2000US-0179065 2000US-0180628 2001WO-US01354

2000US-0209467. 2000US-0214886. 2000US-0215135.

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2000US-0229287. 2000US-0229343. 2000US-0229344. 2000US-0229345. 2000US-0229345. 2000US-0229509. 2000US-0229513. 2000US-0230437. 2000US-0231242.

2000US-0227009 2000US-0228924 2000US-0226868 2000US-0227182 2000US-0225270. 2000US-0225447. 2000US-0225757. 2000US-0225758.

2000US-0225759 2000US-0226279

2000US-0224519. 2000US-0225213. 2000US-0225214. 2000US-0225266. 2000US-0225267. 2000US-0225268.

2000US-0216880. 2000US-0217487. 2000US-0217496. 2000US-0218290. 2000US-0220963.

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Human immune/haematopoietic antigen genomic sequence
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09-AUG-2001
                                  WO200157182-A2
                                                                                                        cytostatic; gene therapy; vaccine; metastasis;
                                                                                                                               Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;
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2000US-0251990.
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Nucleic acids encoding useful for preventing, metastasis -Disclosure; SEQ ID NO 25453; 3071pp + Sequence Listing; English

human immune/hematopoietic antigen polypeptides, diagnosing and/or treating cancers and

amino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic activity, and can be used in gene therapy and vaccine production. (I) proteins and polynucleotides may be used in the prevention, diagnosis and treatment of diseases associated with inappropriate (I) expression. For example, they may be used to treat disorders associated with decreased expression by rectifying mutations or deletions in a patient's genome that affect the activity of (I) by expressing inactive proteins or to supplement the patients own production of (I). Additionally, (I) polynucleotides may be used to produce the secreted (I), by inserting the nucleic acids into a host cell and culturing the cell to express the protein. (I) proteins and polynucleotides may be used to prevent, diagnose and treat immune/haematopoietic related diseases, especially cancers and cancer metastases of haematopoietic antigen genomic sequences from the present invention. AAK54942 to AAK54950 and AAM82169 represent sequences used in the exemplification of the present invention to AAK64702 encode the human immune/haematopoietic antigen ppoietic antigen (I)
(I) have cytostatic

Sequence 12638 BP; 2397 A; 3383 C; 3994 <u>ი</u> 2864 T; 0 other;

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                                 gacatcttcgagaacaagacggacgtggtcctcatcctggagctggtctctggcggggag
                                             gacgtctatgagaaccgcaccgacgtggtgcacatccttgagctagtgtctggaggagag
                                                                               atcgagcgggaggtgaacatcctgcgggagatccggcaccccaacatcatcaccctgcac
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RESULT 1
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                      Query Match
Best Local
                                                                                                                                                                                         The present invention relates to bone marrow expressed polynucleotides and proteins. These sequences can be used in the treatment of inflammatory conditions (eg arthritis, Crohn's disease), cancer, centra and peripheral nervous system diseases and neuropathles, such as Alzheimer's, Parkinson's and Huntington's diseases, spinal cord disorders, head trauma, cerebrovascular diseases, myeloid and lymphoid cell disorders, platelet disorders, eell disorders, bone degenerative disorders, autoimmune disorders, for example multiple sclerosis, diabetes and arthritis, viral and bacterial infections, allergies and blood coagulation disorders. The present sequence is a DN allergies and blood coagulation disorders.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     New bone marrow-expressed nucleic acids and polypeptides, useful for diagnosis, treatment of inflammatory, autoimmune, neurological, cancer and increasing hematopoiesis, stem cell survival and bone growth and
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                31-MAR-2000; 2000US-0540217
23-AUG-2000; 2000US-0649167
23-AUG-2000; 2000US-0649267
30-NOV-2000; 2000US-250583P
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antiparkinsonian; neuroprotective; nootropic; haemostatic; osteopathic;
antiloer; fungicide; antiAdiabetic; antiasthmatic; antiallergic;
immunostimulant; analgesic; cerebroprotective; antianaemic; infection;
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                                                                                                                                                                          allergies and blood of the invention.
                                                                                                                         Sequence
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      and increasing hematopoiesis,
remodeling -
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DB; ABB12364.
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isorder; autoimmune disorder; inflammation;
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                           96
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                         . 88;
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                         Score 228.
Pred. No.
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  Mismatches
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                         .8; DB
3e-55;
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  Indels
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                                                Length
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1;
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Gaps
                                                                                                                                                                                                                                                                                                                                                                        central
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23-AUG-2000;
23-AUG-2000;
30-NOV-2000;
The present invention relates to bone marrow expressed polynucleotides and proteins. These sequences can be used in the treatment of inflammatory conditions (eg arthritis, Crohn's disease), cancer, central and peripheral nervous system diseases and neuropathies, such as Alzheimer's, Parkinson's and Huntington's diseases, spinal cord disorders, head trauma, cerebrovascular diseases, myeloid and lymphoid cell disorders, platelet disorders, stem cell disorders, bone degenerative disorders, autoimmune disorders, for example multiple
                                                                                                                                                                                                                         WPI;
                                                                                                                                                                                                                                                     Tang
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   antiinflammatory; antibacterial; immunosuppressive; vasotropic; cancer; antipakinsonian; neuroprotective; notropic; haemostatic; osteopathic; antialider; fungicide; antidiabetic; antiasthmatic; antiallergic; immunostimulant; analgesic; cerebroprotective; antianaemic; infection;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Human; bone marrow; cytostatic; a antiinflammatory; antibacterial; antiparkinsonian; neuroprotective
                                                                                                                                                                New bone marrow-expressed nucleic acids and polypeptides, useful diagnosis, treatment of inflammatory, autoimmune, neurological, can increasing hematopolesis, stem cell survival and bone growth
                                                                                                                           Claim 1; Page
                                                                                                                                                                                                                                                                              (HYSE-)
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                                                                                                                                                                                                                                                                                                                                                                                                        11-OCT-2001.
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                                                                                                                                                                                                                                                     Boyle
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inflammation; allergy; ds
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                                                                       cancer, central
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                                                                                                                                                                                                                                                                                                                                                                                                         sclerosis, diabetes and arthritis, viral and bacterial infections, allergies and blood coagulation disorders. The present sequence is a DNA of the invention.
                                                                                                                                                                                                                                                                                                                                                                                Sequence 1505 BP; 417 A; 355 C; 399 G; 334 T; 0 other;
                                                                                                                                                          778 cacccctggatc 789
|||||||||||
263 cacccctggatc 274
                                                                                                                                                                                                            203 attcggaagcttctgggttaagagacccggaaacgggtcacaatccaagaggctctcaga
                                                                                                                         718 attcggaagcttctggttaaagagacccggaaacggctcaccaatccaagaggctctcaga 777
                                                                                                                                                                                                                                                                                                                  1; Gaps
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